### PATENT COOPERATION TREATY

## **PCT**

REC'D	12	JUL	2004

WIPO PCT

### INTERNATIONAL PRELIMINARY EXAMINATION REPORT

(PCT Artcle 36 and Rule 70)

Applicant's or agent's file reference 03PCP0006	FOR FURTHER ACTION	SeeNotificationofTransm Examination Report (Form	ittalofInternationalPreliminary				
International application No. PCT/KR2003/000631	International filing date(day/mode) 28 MARCH 2003 (28.03.)	nth/year) Priority d	ate (day/month/year) CH 2002 (29.03.2002)				
		<del></del>	CH 2002 (29.03.2002)				
International Patent Classification (IPC) or national classification and IPC  IPC7 C07H 21/00							
Applicant  CREAGENE INC. et al							
<ol> <li>This international preliminary examination report has been prepared by this International Preliminary Examining Authority and is transmitted to the applicant according to Article 36.</li> <li>This REPORT consists of a total of4sheets, including this cover sheet.</li> <li>This report is also accompanied by ANNEXES, i.e., sheets of the description, claims and/or drawings which have been amended and are the basis for this report and/or sheets containing rectifications made before this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions under the PCT).</li> </ol>							
These annexes consist of a total	ofsheets.						
3. This report contains indications relating to the following items:  I X Basis of the report  II Priority  III Non-establishment of opinion with regard to novelty, inventive step and industrial applicability  IV Lack of unity of invention  V X Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement  VI Certain documents cited  VII Certain defects in the international application  VIII Certain observations on the international application							
Date of submission of the demand	Date o	f completion of this report					
24 OCTOBER 2003 (24.10.2003)		22 JUNE 2004 (22.06.20	004)				
Name and mailing address of the IPEA/KR		rized officer	200				
Korean Intellectual Property Office 920 Dunsan-dong, Seo-gu, Daejeon 302-701, Republic of Korea		AHN, Kyu Jeong					
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### INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International aplication No.
PCT/KR2003/000631

I. Bas	is of the report				
1. With	a regard to the elements of the international application:*				
X	the international application as originally filed				
	the description:				
-	pages	, as originally filed			
	pages	, filed with the demand			
İ	pages, filed with the letter of				
	the claims:				
[	pages	, as originally filed			
1	pages, as amended (together with pages	M1 1 1.0			
[	pages, filed with the letter of	, med with the demand			
	the drawings:				
	pages	, as originally filed			
	pages	, filed with the demand			
	pages, filed with the letter of				
	the sequence listing part of the description:				
	pagespages	, as originally filed , filed with the demand			
	pages, filed with the letter of	, med with the demand			
uie	the language of a translation furnished for the purposes of international search (under Rule the language of publication of the international application (under Rule 48.3(b)).	glish which is 23.1(b)).			
	the language of the translation furnished for the purposes of international preliminary exact or 55.3).				
pre	th regard to any nucleotide and/or amino acid sequence disclosed in the international ap- liminary examination was carried out on the basis of the sequence listing:	pplication, the international			
	contained inthe international application in written form.				
X	filed together with the international application in computer readable form.				
	furnished subsequently to this Authority in written form.				
	furnished subsequently to this Authority in computer readable form				
	The statement that the subsequently furnished written sequence listing does not go beyond the disc local international applications as filed has been furnished.  The statement that the information recorded in computer readable form is identical to the written sequence been furnished.				
4.	The amendments have resulted in the cancellation of:	· · · · · · · · · · · · · · · · · · ·			
	the description, pages the claims. Nos.				
5.	the drawings, sheet				
	This report has been established as if (some of) the amendments had not been made, sing go beyond the disclosure as filed, as indicated in the Supplemental Box(Rule 70.2(c)).**	ce they have been considered to			
• Repla in this and 70	cement sheets which have been furnished to the receiving Office in response to an invitation to opinion as "originally filed." and are not annexed to this report since they do not contait 0.17).	under Article 14 are referred to in amendments (Rules 70.16			
** Any re	placement sheet containing such amendments must be referred to under item I and annexed	to this report.			

#### INTERNATIONAL PRELIMINARY EXAMINATION

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# V. Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1. Statement

	Novelty (N)	Claims	1-20	YES
		Claims	None	·NO
	Inventive step (IS)	Claims	1-2, 4, 8, 13, 17	YES
		Claims	3, 5-7, 9-12, 14-16, 18-20	NO
Industria	Industrial applicability (IA)	Claims	1-20	YES
		Claims	None	NO

#### 2. Citations and explanations (Rule 70.7)

The following documents have been considered for the purpose of this report:

D1: US 6361939 B1 (26 March 2002)

D2: J. of Biol. Chem., Vol. 276(21): 17920-17931 (2001)

D3: Gene Accession number AL354813 (16 May 2001)

D4: Gene Accession number AF067844 (08 February 1999)

The present invention relates to polynucleotides highly expressed in subsets of dendritic cells (DCs) and matured DCs, and microarrays comprising the same. These polynucleotides are characterized by the expression in CD11-DCs from peripheral blood, CD1a<sup>+</sup> or CD14<sup>+</sup> DCs generated from CD34<sup>+</sup> progenitor cells isolated from umbilical cord blood.

#### Novelty

D1 and D2 disclose polynucleotides encoding DC-specific proteins associated with DC differentiation and maturation. D2 also discloses the analysis of the expression profile of human CD14+ blood monocytes and their derived DCs using DNA microarrays.

The polynucleotides of SEQ ID NOs: 4 and 6 of this invention was published in the NCBI database (D3; D4) before the priority date (29 March 2002) as a part of the genomic sequence. Said polynucleotides are disclosed in D3 and D4 merely as a part of the genomic sequence without any function. In addition, the DC-specific expression of said polynucleotides is not mentioned in D3 and D4.

None of prior art discloses the DC-specific polynucleotides consisting of SEQ ID No: 1, 3, 4, 5, or 6.

Thus, the novelty of this invention can be acknowledged (PCT Article 33(2)).

(Continued on Supplemental Sheet.)

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Supplemental Box

(To be used when the space in any of the preceding boxes is not sufficient)

Continuation of:

BoxV. 2.

Inventive Step

Claims 3, 5-7, 9-12, 14-16 and 18-20 relate to a method for detecting a dendritic cell or subset of DCs, comprising the hybridization of DNA obtained from cells with a DC-specific nucleotide sequence; a method for detecting a maturation stage of DC; and microarrays for these methods. For these methods, the present invention uses genes for Ig superfamily protein, DC-LAMP, interferon regulatory factor 4 (IRF4), and RNase A, which are already disclosed in D1 and D2.

D1 discloses a kit comprising polynucleotide encoding LAMP-like family member, and D2 discloses a method for detecting genes associated in DC differentiation and maturation using microarray.

It appears obvious to a person skilled in the art to arrive at the present claims 3, 5-7, 9-12, 14-16 and 18-20 without the exercise of inventive skill. Therefore, the subject matter of claims 3, 5-7, 9-12, 14-16 and 18-20 lack an inventive step under PCT Article 33(3).

**Industrial Applicability** 

The subject matter of claims 1-20 are considered to be industrially applicable (PCT Article 33(4)).

**New Citation** 

- 1. Gene Accession number AL354813 (16 May 2001)
- 2. Gene Accession number AF067844 (08 February 1999)